Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

Claim 1. (Currently amended) A pharmaceutical formulation for the controlled release of a therapeutic peptide or a salt of the peptide, said formulation comprising: thereof,

a peptide comprising which peptide has the sequence (SEQ ID NO: 7)

pyro Glu-His-Trp-Ser-Xaa¹-Gly-Xaa²-Xaa³-Pro-Gly-NH₂, or a salt thereof, wherein Xaa¹ is His or Tyr,

Xaa² is Trp or Leu, and

Xaa³ is Tyr or Arg,

provided that when Xaa¹ is Tyr and Xaa² is Leu, then Xaa³ is not Arg,

and which formulation further comprises a pharmaceutically acceptable biodegradable polymer.

Claim 2. (Currently amended) The pharmaceutical <u>formulation</u> eomposition according to Claim 1, wherein the peptide <u>comprises</u> is (SEQ ID NO: 6)

pyroGlu-His-Trp-Ser-His-Gly-Trp-Tyr-Pro-Gly-NH₂.

Claim 3. (Currently amended) The <u>pharmaceutical</u> formulation according to Claim 1, wherein the polymer <u>comprises</u> is a polymer of a hydroxyl derivative of a carboxylic acid, <u>a</u> <u>polymer of an amino derivative of a carboxylic acid</u>, or a copolymer of <u>such the carboxylic acid</u> derivatives.

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Claim 4. (Currently amended) The <u>pharmaceutical</u> formulation according to Claim 3, wherein the polymer <u>comprises</u> is a polymer of glycolic acid, a polymer of lactic acid, or a copolymer of <u>glycolic and lactic acid</u> such derivatives.

Claim 5. (Currently amended) The <u>pharmaceutical</u> formulation according to Claim 1, wherein the peptide is microencapsulated by the polymer.

Claim 6. (Currently amended) A method for treating a human pathology comprising administering the treatment of a human medical condition, which method comprises the administration to an individual in need of said such treatment of a therapeutically effective amount of the pharmaceutical formulation a controlled release formulation of a peptide according to Claim 1 any of the preceding claims.

Claim 7. (Currently amended) A-The pharmaceutical formulation according to claim 1 for treatment of, or for protection against, <u>a</u> disorder of bone growth or disorder of prostate growth.

Claim 8. (Currently amended) The use of peptide or salt as defined in claim 1, together with a pharmaceutically acceptable biodegradable polymer, for the preparation of A method for preparing a controlled release medicament for the treatment of, or protection against, a disorder of bone growth or prostate growth comprising:

providing a peptide comprising (SEQ ID NO: 7)

pyro Glu-His-Trp-Ser-Xaa¹-Gly-Xaa²-Xaa³-Pro-Gly-NH₂, or a salt thereof,

wherein Xaa¹ is His or Tyr,

Xaa² is Trp or Leu, and

Xaa³ is Tyr or Arg,

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provided that when Xaa¹ is Tyr and Xaa² is Leu, then Xaa³ is not Arg, and

incorporating the peptide or salt into a matrix of a pharmaceutically acceptable biodegradable polymer or encapsulating the peptide or salt with a pharmaceutically acceptable biodegradable polymer.

Claim 9. (Currently amended) A use The method according to claim 8, wherein said polymer is comprises:

- (a) a polymer of a hydroxy derivative of a carboxylic acid, a polymer of an amino derivative of a carboxylic acid, or a copolymer of the carboxylic acid such derivatives, or
- (b) a polymer of glycolic acid, a polymer of lactic acid, or a copolymer of lactic and glycolic acids.

Claim 10. (Currently amended) A use The method according to claim 8, wherein said disorder is selected from the group consisting of age-related osteoporosis, osteoporosis associated with post-menopausal hormone status, primary and secondary hyperparathyroidism, disuse osteoporosis, diabetes-related osteoporosis, and glucocorticoid-related osteoporosis, benign prostatic hyperplasia and prostate cancer.

Claim 11. (Currently amended) A-The pharmaceutical formulation according to claim 7, for the treatment of, or protection against, wherein said disorder is selected from the group consisting of selected form age-related osteoporosis, osteoporosis associated with post-menopausal hormone status, primary and secondary hyperparathyroidism, disuse osteoporosis, diabetes-related osteoporosis, and glucocorticoid-related osteoporosis, benign prostatie hyperplasia and prostate cancer.

Claim 12. (Currently amended) A The method according to claim 6, wherein the human pathology comprises for treating or protecting against a human disorder of bone growth or of

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prostate growth, which method comprises the administration to an individual in need of such treatment or protection of a therapeutically effective amount on a formulation according to claim 1.

Claim 13. (Currently amended) A The method according to claim 12, wherein said disorder is selected from the group consisting of age-related osteoporosis, osteoporosis associated with post-menopausal hormone status, primary and secondary hyperparathyroidism, disuse osteoporosis, diabetes-related osteoporosis, and glucocorticoid-related osteoporosis, benign prostatic hyperplasia and prostate cancer.